Bioterrorism Preparedness -Laboratory Analysis

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Bioterrorism Preparedness -Laboratory Analysis

An account from the "real world" of the clinical microbiology laboratory

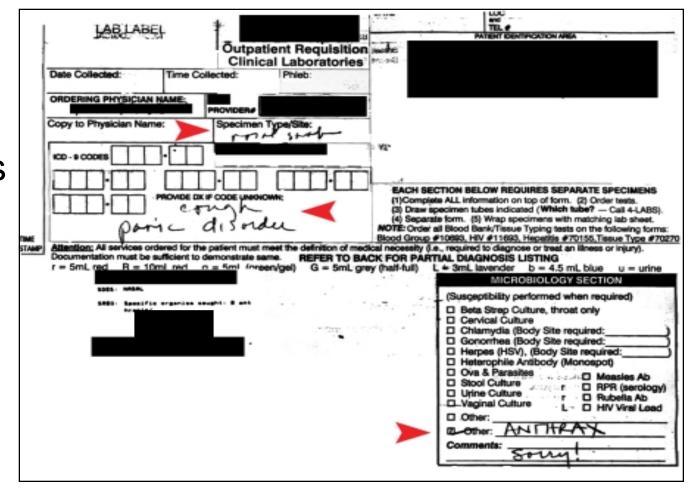


Clinical Laboratories -The Need for Preparation

- Agents likely to be used by terrorists
 - Unfamiliar, rarely encountered organisms
 - Potential for misidentification, mishandling of specimens, laboratory acquired infection
- Public health agency-sponsored training in the Northeast began in 1999
- Laboratory Response Network (LRN)
- Were we prepared in the autumn of 2001?

Autumn, 2001 - Anthrax!

- Wake-up call for clinical microbiologists
- Expect the unexpected
- Preparedness is an absolute necessity



LRN Level A Lab Preparedness

- Level A laboratory functions
 - Rule out / refer
 - Ship suspicious infectious agents to higher level labs for further study
- Level A laboratory activities
 - Formulate laboratory procedures
 - Train staff
 - Biosafety concerns
- Assistance from public health agencies

Activities of Clinical Micro Labs

- "Average" Labs
 - Microscopic examination of specimens
 - Culture of specimens and isolation of many bacterial and fungal pathogens
 - Identification and susceptibility testing
- "Advanced" Labs
 - Viruses (culture, direct detection)
 - Mycobacteria (culture, susceptibility)
 - Certain fungi (culture and identification)
 - Molecular testing

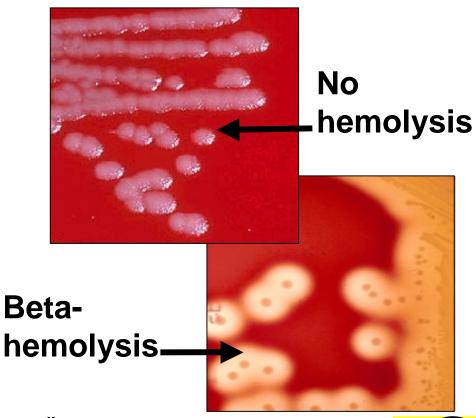
 Gram stain* of CSF, positive blood culture or wound culture shows large grampositive rods



*Gram stain: Differential stain, not specific, but can be extremely helpful



Culture on blood agar*.
 Examine for characteristic colony morphology and lack of beta-hemolysis



*Agents of anthrax and plague are "easy" to grow. Agents of tularemia, brucellosis are harder to recover, may require special media



 Perform identification tests. For ?B. anthracis, perform motility test*

Growth throughout medium (motile)

Growth only near original inoculation stab (non-motile)

*Minimal rule out tests (minimal manipulation of potentially dangerous cultures) are recommended for Level A labs



- Ruled in?
 - Bacillus species with characteristic colony morphology, non-hemolytic, non-motile
- REFER
 - Contact Level B lab
 - Ship suspect isolate



Level A Lab Preparedness - Where Are We Now?

- Bigger seems to be "better"
 - Wider variety of pathogens encountered; personnel experienced in working with infrequently isolated agents
 - More and/or better biosafety equipment
 - Institutional support for needed resources is more likely in larger hospitals
- Small labs can still have successful preparedness programs

Level A Lab Preparedness





Anthrax





Brucellosis





Plague



Botulism-Specimen processing/ shipping only





Tularemia



Smallpox, VHF-More guidance needed for Level A labs



Environmental testing for *B. anthracis spores*

Clinical Lab Preparedness – Next Steps

- Extend training (category B agents)
- Enhance communication/cooperation with higher level public health labs
 - NLS
- Dissemination of some Level B procedures to select Level A labs
 - ?Rapid, specific tests/reagents
 - ?BSL3 activities in select labs
 - ?Surge capacity

Level A Clinical Microbiology Laboratories

- Can be instrumental in early recognition
- Must be trained, alert and vigilant
- Form partnerships with public health labs for BT preparedness assistance, BT response plans, and overall improvement of the public health system